

KEYED INTERVERTEBRAL DOWEL

This application is a Continuation of U.S. patent application Serial No.09/327,982, which was filed June 8, 1999, and is hereby incorporated herein by reference.

5 BACKGROUND OF THE INVENTION

1. Technical Field

 The present disclosure relates to an intervertebral implant for spinal fusion and more particularly, to an intervertebral dowel having at least two radially extending tabs for securing the dowel within a receiving bed formed in the
10 intervertebral space.

2. Background of Related Art

 The spine is a flexible column formed of a series of bone called vertebrae. The vertebrae are hollow and piled one upon the other, forming a strong hollow column for support of the cranium and trunk. The hollow core of the spine
15 houses and protects the nerves of the spinal cord. The different vertebrae are connected together by means of articular processes and intervertebral, fibro-cartilages. In general, a vertebral body is made of a cortical shell enclosing a cancellous (spongy) bone core. The portion of the cortical bone shell facing the surface of the disk is the endplate.

20 The intervertebral fibro-cartilages are also known as intervertebral disks and are made of a fibrous ring filled with pulpy material. The disks function as spinal shock absorbers and also cooperate with synovial joints to facilitate movement and maintain flexibility of the spine. When one or more disks degenerate through trauma,

spondylolisthesis or other pathologies, nerves passing near the affected area may be compressed and are consequently irritated. The result may be chronic and/or debilitating back pain. Various methods and apparatus, both surgical and non-surgical, have been designed to relieve such back pain.

5 One method designed to relieve such back pain is interbody spinal fusion. Typically, interbody spinal fusion involves distracting adjoining vertebrae of the spine so that the nerve root canal sizes are increased and nerve irritation is eliminated or reduced. In order to maintain the adjoining vertebrae in a distracted state, at least one intervertebral implant is inserted into a receiving bed formed between
10 the vertebrae. The implant is positioned to engage the adjoining vertebrae to maintain the vertebrae at a fixed degree of distraction.

 Preferably, the implant should stabilize the intervertebral space and become fused to adjacent vertebrae in order to prevent the implant and adjacent vertebrae from moving. The implant must also provide spinal load support between the
15 vertebrae. Further, during the time it takes for fusion, i.e. biological fixation of the vertebrae, to be completed, the implant should have enough structural integrity to maintain the space without substantial degradation or deformation of the implant. The implant should also have sufficient stability to remain in place prior to actual completion of bone ingrowth fusion. The implant should include structure which
20 maintains the implant in position between the vertebrae while bone ingrowth is occurring. To facilitate rapid bone growth, and thus quick fusion, the implant may

include or be provided with a bone growth supporting material. Obviously, the material from which the implant is constructed should be a biocompatible material and, preferably, interact biologically with the body's own naturally occurring tissues.

A variety of different types of intervertebral implants have been developed to perform this function including spinal fusion cages, threaded bone dowels and stepped bone dowels. An exemplary implant is disclosed in U.S. Patent Application filed on even date herewith, under Certificate of Express Mail Label No. EL260888076US, and entitled "Ramp-Shaped Intervertebral Implant", the entire disclosure of which is incorporated by reference herein.

Common deficiencies in some of the prior art implants may include expulsion of the implant from between adjacent vertebrae, difficulty in inserting the implant into position, and/or lack of ability to allow incorporation of implant into the body. Also, in some prior art spinal fusion methods utilizing implants, the vertebrae may need to be distracted to a large extent in order to position the implant between the vertebrae.

Accordingly, a need exists for an improved intervertebral implant which is configured to prevent the likelihood of expulsion or retropulsion during normal patient activity, provide ease of insertion and include structure to facilitate incorporation of the implant into the body. Furthermore, need exists for an improved intervertebral implant which can be inserted between vertebrae without excessive distraction of the vertebrae and a method of installing such an implant.

SUMMARY

In accordance with the present disclosure, an intervertebral implant having tabbed securing structure is provided. The intervertebral implant includes a substantially cylindrical body portion and at least one pair of radially extending tabs that are configured to engage vertebral bodies.

By engaging the vertebrae, the tabs reduce the likelihood that expulsion or retropulsion might occur. This is particularly significant in that where an implant is pushed out of place, damage to vital structures including neural (the spinal cord and existing nerve roots) and vascular (the aorta and inferior vena cava) can occur resulting in possible injury or death. Additionally, the tabs assist in preventing migration of the implant due to rotation of the adjacent vertebrae.

The tabs may take the form of various shape and constructions, such as, for example, smooth rounded, wedge shaped, cam shaped, toothed, or threaded, etc. In alternate embodiments, two diametrically opposed pairs of tabs are provided on the cylindrical body portion. In various embodiments, a throughbore or a plurality of throughbores extend from a top surface of the implant to the bottom surface of the implant providing a space for boney bridging to occur between the vertebrae which are intended to be fused. The throughbore(s) is dimensioned to receive growth factors or other grafting materials to stimulate bone healing. The pairs of tabs may be provided adjacent the opening of the throughbore or may be offset 90° from the openings of the throughbore. In one embodiment of an intervertebral implant, the cylindrical body portion is tapered.

In an alternate embodiment, the implant has an abbreviated body portion and does not include a throughbore.

In another embodiment, the tabs are formed by inserting a cortical plug through the throughbore. Preferably, the cylindrical body portion includes a slot
5 formed in one end thereof for receipt of an insertion tool and a bore extending between the slot and into the throughbore for facilitating insertion and facilitating injection into the throughbore of any desirable material, such as, for example, bone growth stimulants, autograft, allograft, demineralized bone matrix, or other bone grafting materials.

10 Further, alternate embodiments may include body portions having shapes other than cylindrical, such as, those having rectangular, oval, multi-sided, etc., cross-sections.

In a preferred embodiment, the implant is formed from a cortical ring allograft cut from the diaphysis of a long bone. By utilizing bone or bone-derived
15 materials as the implant material, the implant has the added advantage of facilitating incorporation of the implant into the body. The implant can be formed by milling the top and bottom surfaces of a cortical ring to form the substantially cylindrical body portion and a pair of radially extending wings. The implant is further milled such that the radially extending wings are formed into tabs each of which is spaced a
20 predetermined distance from the end of the cylindrical body portion. Additionally, each tab may be milled so as to form the desired camming, wedge, threaded, etc.

least one pair of diametrically opposite radially extending tabs extending from the cylindrical body portion and inserting the implant between adjacent vertebrae such that the tabs are in alignment with the space defined between adjacent vertebrae. The method further includes positioning the implant such that the tabs are within the enlarged areas of the stepped bore and rotating the implant such that the tabs enter the enlarged or stepped area of the bore. This provides a greater ease of insertion over other styles of implants, such as, for example, threaded implants.

BRIEF DESCRIPTION OF THE DRAWINGS

Various preferred embodiments are described herein with reference to the drawings wherein:

FIG. 1 is a perspective view of one embodiment of the presently disclosed intervertebral implant;

FIG. 2 is a side view of the intervertebral implant shown in FIG. 1;

FIG. 3 is a top view of the intervertebral implant shown in FIG. 1;

FIG. 4 is a front view of the intervertebral implant shown in FIG. 1;

FIG. 5 is a side view of a long bone;

FIG. 6 is a perspective view of a ring cut from the long bone shown in FIG. 5;

FIG. 7 is a side view of the ring shown in FIG. 6;

FIG. 8 is a perspective view of the ring after the top surface has been milled;

FIG. 9 is a perspective view of the ring after the bottom surface has been milled;

FIG. 10 is a perspective view of the ring after the side walls have been machined;

5 FIG. 11 is a perspective view of the ring after the radially extending wings have been machined to form tabs;

FIG. 12 is a an end view of the vertebral space with a stepped hole drilled therein;

FIG. 13 is a side view of the vertebral space shown in FIG. 12;

10 FIG. 14 is an end view of the vertebral space of FIG. 12 with one embodiment of the presently disclosed intervertebral implant inserted therein;

FIG. 15 is a perspective view similar to FIG. 14 with the intervertebral implant rotated 90°;

15 FIG. 16 is a side view of the intervertebral space similar to FIG. 13 with the intervertebral implant inserted and rotated 90°;

FIG. 17 is a perspective view of another embodiment of the presently disclosed intervertebral implant;

FIG. 18 is a side view of the intervertebral implant shown in FIG. 17;

FIG. 19 is a top view of the intervertebral implant shown in FIG. 17;

20 FIG. 20 is a front view of the intervertebral implant shown in FIG. 17;

FIG. 21 is a perspective view of another embodiment of the presently disclosed intervertebral implant;

FIG. 22 is a side view of the intervertebral implant shown in FIG. 21;

FIG. 23 is a top view of the intervertebral implant shown in FIG. 21;

5 FIG. 24 is a front view of the intervertebral implant shown in FIG. 21;

FIG. 25 is a perspective view of another embodiment of the presently disclosed intervertebral implant;

FIG. 26 is a side view of the intervertebral implant shown in FIG. 25;

FIG. 27 is a top view of the intervertebral implant shown in FIG. 25;

10 FIG. 28 is a front view of the intervertebral implant shown in FIG. 25;

FIG. 29 is a perspective view of another embodiment of the presently disclosed intervertebral implant;

FIG. 30 is a side view of intervertebral implant shown in FIG. 29;

FIG. 31 is a top view of the intervertebral implant shown in FIG. 29;

15 FIG. 32 is a front view of the intervertebral implant shown in FIG. 29;

FIG. 33 is a perspective view of another embodiment of the presently disclosed intervertebral implant;

FIG. 34 is a side view of the intervertebral implant shown in FIG. 33;

FIG. 35 is a top view of the intervertebral implant shown in FIG. 33;

20 FIG. 36 is a front view of the intervertebral implant shown in FIG. 33;

FIG. 37 is a perspective view of another embodiment of the presently disclosed intervertebral implant;

FIG. 38 is a side view of the intervertebral implant shown in FIG. 37;

FIG. 39 is top view of the intervertebral implant shown in FIG. 37;

5 FIG. 40 is a front view of the intervertebral implant shown in FIG. 37;

FIG. 41 is a perspective view of another embodiment of the presently disclosed intervertebral implant;

FIG. 42 is a side view of the intervertebral implant shown in FIG. 41;

FIG. 43 is a top view of the intervertebral implant shown in FIG. 41;

10 FIG. 44 is a front view of the intervertebral implant shown in FIG. 41;

FIG. 45 is a perspective view of another embodiment of the presently disclosed intervertebral implant;

FIG. 46 is a side view of the intervertebral implant shown in FIG. 45;

FIG. 47 is a top view of the intervertebral implant shown in FIG. 45;

15 FIG. 48 is a front view of the intervertebral implant shown in FIG. 45;

FIG. 49 is a perspective view of another embodiment of the presently disclosed intervertebral implant;

FIG. 50 is a side view of the intervertebral implant shown in FIG. 49;

FIG. 51 is a top view of the intervertebral implant shown in FIG. 49;

FIG. 52 is a front view of the intervertebral implant shown in FIG. 49;

FIG. 53 is a perspective view of another embodiment of the presently disclosed intervertebral implant;

FIG. 54 is a side view of the intervertebral implant shown in FIG. 53;

5 FIG. 55 is a top view of the intervertebral implant shown in FIG. 53;

FIG. 56 is a front view of the intervertebral implant shown in FIG. 53;

FIG. 57 is a perspective view of another embodiment of the presently disclosed intervertebral implant;

FIG. 58 is a side view of the intervertebral implant shown in FIG. 57;

FIG. 59 is a top view of the intervertebral implant shown in FIG. 57;

FIG. 60 is a front view of the intervertebral implant shown in FIG. 57;

FIG. 61 is a perspective view of another embodiment of the presently disclosed intervertebral implant body portion with a rectangular cross-section;

FIG. 62 is a perspective view of another embodiment of the presently
15 disclosed intervertebral implant body portion with an oval cross-section; and

FIG. 63 is a perspective view of another embodiment of the presently disclosed intervertebral implant body portion with a multi-sided cross-section.

DETAILED DESCRIPTION OF PREFERRED EMBODIMENTS

Preferred embodiments of the presently disclosed intervertebral implant
20 will now be described in detail with reference to the drawings, in which like reference
numerals designate identical or corresponding elements in each of the several views.

portion 12 has a first end 18 and a second end 20. Tab 14 has first and second engaging or retaining surfaces 22a and 22b which are stepped or longitudinally spaced a predetermined distance from first end 18 and second end 20, respectively. Similarly, tab 16 has a pair of retaining surfaces 24a and 24b which are similarly stepped or longitudinally spaced from a first end 18 and second end 20 respectively. Retaining surfaces 22a, 22b and 24a, 24b are configured to engage a portion of adjacent vertebrae when installed therebetween.

As shown, tabs 14 and 16 extend only along a limited extent of the circumference of a cylindrical body portion 12. Preferably, tabs 14 and 16 are radially spaced 180° apart. Tab 14 includes a rounded side surface 26 and tab 16 includes a rounded side surface 28.

As shown, implant 10 includes a throughbore 30 which has a longitudinal axis substantially perpendicular to the longitudinal axis of implant 10. Further, implant 10 may be provided with perforations instead of, or in addition to, throughbore 30. Where implant 10 is formed of bone, the perforations assist in facilitating biological attachment and eventual incorporation of the implant into adjacent vertebrae.

Implant 10 further includes an installation slot 32 machined or milled in first end 18. A second bore 34 extends between slot 32 and throughbore 30. Second bore 34 is provided for mating of the implant with an insertion tool. Throughbore 30 is dimensioned to receive bone particles and/or biocompatible osteoinductive or

osteochonductive material. These materials may include cancellous bone, cancellous bone particles, ceramics, polymers, composites, BMP, etc.

Intervertebral implant 10 can be constructed from a broad range of biocompatible materials such as, for example, surgical stainless steel, titanium, ceramic, hydroxyapatite, polymer, carbon fiber, tantalum, etc. Preferably, implant 10 is constructed from a human and/or animal cadaver bone. Intervertebral implant 10, appropriately sized, can be used in cervical, thoracic and lumbar spinal fusion procedures. For cervical spinal fusion procedures, in which implants are typically between 8 to 15 mm in length and 10 to 14 mm in diameter, bone is preferably obtained from the fibula, radius, ulna or humerus bones. For thoracic and lumbar spinal fusion procedures in which implants are typically 10 to 30 mm in length and/or diameter and about 10 to 14 mm in height, bone is preferably obtained from the humerus, femur or tibia. The sources of cortical bone for the bone-derived implant are preferably allogenic but also include xenogenic sources such as bovine and porcine bone.

Additionally, the bone may be subjected to penetration with osteogenic or demineralization agents during manufacture of the implant.

Alternatively, as discussed above, intervertebral implant 10 can be molded or machined from other biocompatible materials including composites made of bone as discussed in U.S. Patent No. 5,899,939 to Boyce et al., the entire disclosure of which is incorporated by reference herein.

biocompatible material. As such, other known processes, such as molding techniques may be used to manufacture the implant.

Installation of implant 10 between a pair of adjacent vertebrae will now be described. Referring to Figs. 12-16 and initially to Figs. 12-13, there is illustrated a pair of adjacent vertebrae X and Y defining intervertebral space Z therebetween. The endplate is stronger bone than is the cancellous core. Thus, cuts in the vertebral bodies permit the tabs of the implant to extend past the endplate and into the softer bone beneath. A camming approach for some of the following disclosed embodiments of the implant tabs allows the cancellous bone to be compressed against the implant thereby providing additional frictional resistance against implant movement. A drill or other known devices and methods are utilized to form a stepped hole or bore E between the adjacent vertebrae preferably by milling or machining. Examples of such devices and procedures are disclosed in U.S. Patent No. 5,445,639, the entire disclosure of which is incorporated by reference herein. Stepped hole E preferably has narrow diameter portion F adjacent the outer surface of the vertebrae and enlarged portion G interior to the vertebrae. In preparation for use, intervertebral implant 10 may be demineralized as discussed hereinabove and mounted on suitable installation devices.

Referring now to Fig. 14, once installed on an insertion device, intervertebral implant 10 is inserted between vertebrae X and Y such that tabs 14 and 16 are aligned with the intervertebral space Z. Intervertebral implant 10 is inserted into the drilled hole a sufficient distance such that tabs 14 and 16 align with the

engaging surfaces 62a and 64a for engaging an edge of stepped bore in a drilled vertebrae. Similarly, second tabs 66 and 68 also include engaging surfaces 66a and 68a for engaging an interior of a bore drilled in bone or vertebrae. Similar to that disclosed with regard to implant 10, first tabs 62 and 64 as well as second tabs 66 and 68 may have a generally rounded profile.

Intervertebral implant 50 is formed in the manner disclosed above with respect to implant 10 and is similarly installed in a stepped bore drilled in adjacent vertebrae. The stepped bore may have only a single enlarged area or may include two separate enlarged areas to accommodate the first and second tabs as the intervertebral implant is rotated into place.

Referring now to FIGS. 21-24, there is disclosed another alternate embodiment of an intervertebral implant similar to that of implant 50. Intervertebral implant 80 includes a generally cylindrical body portion 82 having a throughbore 84 formed therethrough. An installation slot 86 is provided along with a bore 88 extending between installation slot 86 and an interior of throughbore 84. Implant 80 includes a pair of radially extending first tabs 90 and 92 as well as a pair of radially extending second tabs 94, 96. In contrast to implant 50, first tabs 90, 92 and second tabs 94, 96 are formed on cylindrical body portion such that they are generally perpendicular to slot 86 and are adjacent to throughbore 84.

In the presently disclosed embodiments where the tabs are adjacent to the throughbore, a different method of forming the implant from bone is necessary.

Referring now to FIGS. 25-28, there is disclosed another embodiment of an intervertebral implant which includes specific wedging structure to prevent the implant from moving longitudinally within a bore. Implant 100 generally includes a cylindrical body portion 102 having a throughbore 104 formed therein. Similar to previous embodiments, implant 100 is provided with an installation slot 106 and a bore 108 extending between installation slot 106 and throughbore 104. Implant 10 also includes a pair of radially extending first anterior tabs 110, 112 and a pair of radially extending second tabs 114, 116. As shown, first tabs 110 and 112 have curved wedge surfaces 118, 120. Similarly, second tabs 114 and 116 also include curved wedge surfaces 122 and 124. Wedge surfaces 118 and 120 of first tabs 110 and 112 curve away from a first end 126 of implant 10 and wedge surfaces 122, 124 of second tabs 114 and 116 curve away from a second end 128 of implant 100. The provision of wedge surfaces on the tabs provides a range of camming contact with the interior of a stepped bore drilled in adjacent vertebrae to thereby prevent expulsion of the implant.

20 Referring now to FIGS. 29-32, there is disclosed a further alternate embodiment of an intervertebral implant which includes progressive, radial camming

structure which, upon rotation of the implant, cams the implant into position within a stepped bore. Specifically, intervertebral implant 130 includes a cylindrical body portion 132 having a throughbore 34 formed therethrough. An installation slot 136 may be provided along with a bore 138 extending between installation slot 136 and throughbore 134. Implant 130 additionally includes first tabs 140 and 142 formed adjacent first end 144 and second tabs 146 and 148 formed adjacent a second end 150. As illustrated, first tabs 140 and 142 as well as second tabs 146 and 148 have a generally, progressively curved shape such as a spline shape or one defined by a polynomial-defined curve. Thus, first tabs 140, 142 include progressive camming surfaces 152, 154. Second tabs 146 and 148 include progressive camming surfaces 156 and 158. Implant 130 may be formed in a manner similarly described above with respect to implant 10.

Upon installation of implant 130, between adjacent vertebrae, implant 130 is rotated and progressive camming surfaces 152, 154 and 156, 158 engage walls of the stepped bore in progressive fashion to firmly wedge implant 130 within the stepped bore and prevent any loosening or further rotation or reverse rotation of implant 130 within the stepped bore. The provision of progressive camming surfaces allows for the use of implant 130 in bores which may not have been drilled precisely or to a constant/consistent diameter. Further, as noted above, camming structure on the disclosed implants allows the tabs to compress the spongy bone to gain additional frictional force to secure the implant between the vertebrae.

body portion 192. Implant 190 additionally includes second tabs 202 and 204 positioned adjacent and spaced a distance from second end 206 of cylindrical body portion 192. Implant 190 includes camming structure formed on the first and second tabs which permits rotation of the implant in either direction upon installation.

5 Specifically, first tabs 196 includes opposed inclined camming surfaces 208a and 208b and first tab 198 also includes opposed inclined camming 210a and 210b. Similarly, second tab 202 includes opposed inclined camming surfaces 212a and 212b and second tab 204 includes opposed inclined camming surfaces 214a and 214b. The opposed inclined camming surfaces allow the implant to be rotated in either direction and still
10 achieve a camming function within a stepped bore. As with prior embodiments, first tabs 196 and 198 include bore engaging surfaces 196a and 198a respectively. Similarly, second tabs 202 and 204 include bore engaging surfaces 202a and 204a respectively. Implant 190 may preferably be provided with an installation slot 216 and a bore 218 extending between slot 216 and throughbore 194.

15 Referring now to FIGS. 41-44, there is disclosed a further alternate embodiment of an intervertebral implant. Implant 220 generally includes cylindrical body portion 222 having a throughbore tube 224 defined therein. First tabs 226 and 228 and second tabs 230 and 232 extend radially from cylindrical body portion 222. The first and second tabs of implant 220 include threaded structure which allows the
20 implant to engage precut threads in a stepped bore formed between adjacent vertebrae or to act as teeth to cut into bone and thereby secure implant 220 within a stepped bore

between adjacent vertebrae. Alternatively, the tabs may be grooved but not necessarily threaded. Specifically, first tab 226 includes a threaded surface 234 and first tab 228 includes a threaded surface 236. Similarly, second tab 230 includes a threaded surface 238 and second tab 232 includes a threaded surface 240. It should be noted that the number of threads on any individual tab may differ from the number on an adjacent or diametrically opposed tab. Preferably, an installation slot 242 is provided having a bore 244 extending between slot 242 and into throughbore 224.

Referring now to Figs. 45-48, there is disclosed an asymmetrical embodiment of an intervertebral implant. Implant 250 generally includes a cylindrical body portion 252 having a first end 254 and a second end 256. A throughbore 258 extends through implant. A first tab 260 is provided a predetermined spaced distance from first end 254 and a second tab 262 is provided a predetermined spaced distance from second end 256. As shown, first and second tabs 260, 262 are radially spaced approximately 180°. First and second tabs 260, 262 may be of any of the previously described shapes in the prior embodiments and include respective camming and/or abutment bone engaging surfaces. Additionally, implant 250 may be provided with an installation slot 269 and a bore 266 and be formed in accordance with the previously described methods and of same or similar materials.

Referring now to Figs. 49-52, there is disclosed an intervertebral implant 270 designed to utilize a plug, which may be formed from cortical bone, to form the tabs. Implant 270 generally includes a cylindrical body portion 272 formed in

accordance with the above described method such that the medullary canal provides a throughbore 274 in implant 270. A cortical plug 276 formed by turning on a lathe, milling, or other appropriate machining process. Plug 276 is positioned within throughbore 274 which may be suitably drilled or otherwise prepared to receive plug 276 such that first and second ends 278, 280 of cortical plug 276 extend radially outward from body portion 272. First and second ends 278, 280 thus form tabs which, when installed by the above described method, engage edges of a stepped bore formed in adjacent vertebrae. An installation slot 282 may be formed in an end 284 of body portion and a bore 286 extends between slot 282 and throughbore 274.

Referring now to Figs. 53-56, there is disclosed an alternate embodiment of an intervertebral implant with a substantially shortened body portion. Implant 290 is designed to be provided in various diameters such that two or more implants 290 of differing diameters may be used together to introduce the appropriate lordosis into the spine. Implant 290 generally is similar to the above described implants except that the length of a cylindrical body portion 292 is substantially abbreviated or shortened.

Implant 290 may include any of the previously described versions of tabs and preferably first and second tabs 294, 296. Implant 290 may also include an installation slot 298 and bore 300 extending between slot 298 and end face 302 of body portion 292. However, it is not contemplated that implant 290 have a throughbore and thus implant 290 may be formed from bone extending up to, but not including, the

medullary canal of a long bone. Further, various body portion configurations, such as, for example, tapered, semi-conical, etc. are also envisioned.

Referring now to Figs. 57-60, there is disclosed another embodiment of an intervertebral implant. Implant 310 generally includes a tapered cylindrical body portion 312 having a first end 314 and a second end 316. The diameter of first end 314 is smaller than the diameter of second end 316. Implant 310 may be formed by the disclosed method and include a throughbore 318, an installation slot 320 and a bore 322 extending from slot 320 to throughbore 318. Additionally, implant includes first tabs 324, 326 and second tabs 328, 330.

As best shown in FIGS. 61-63, various body portions other than cylindrical are within the contemplated scope of the present disclosure. These body portions may include a body portion 340, having a rectangular cross-section (FIG. 61), a body portion 350 having an oval cross-section (FIG. 62), a body portion 360 having a multi-sided cross-section (FIG. 63), etc. The embodiments disclosed in FIGS. 61-63 may obviously include structure similar or identical to that provided in previously described embodiments such as, for example, throughbores, installation slots, bore and all the various configurations and orientations of tabs.

It will be understood that various modifications may be made to the embodiments disclosed herein. For example, differing or alternate tab constructions may be provided on a single implant. Additionally, the various configurations may be combined on individual tabs. Therefore, the above description should not be construed

as limiting, but merely as exemplifications of preferred embodiments. Those skilled in the art will envision other modifications within the scope and spirit of the claims appended hereto.